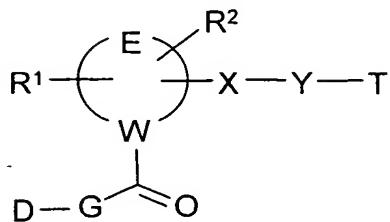


**Patent Claims****1. Compounds of the formula I**

5



10

in which

$R^1$ ,  $R^2$  are each, independently of one another, H, =O, Hal, A, ethynyl, OR<sup>3</sup>, N(R<sup>3</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, N<sub>3</sub>, COOR<sup>3</sup>, CON(R<sup>3</sup>)<sub>2</sub>, [C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-Ar, [C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-Het, [C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-cycloalkyl, -OCOR<sup>3</sup>, -OCON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>COA or NR<sup>3</sup>SO<sub>2</sub>A,

15

$R^1$  and  $R^2$  together are alternatively a bicyclically or spirocyclically bonded 3- to 7-membered carbocyclic or heterocyclic ring having from 0 to 3 N, O and/or S atoms,

20

$R^3$  is H, A, H-C≡C-CH<sub>2</sub>-, CH<sub>3</sub>-C≡C-CH<sub>2</sub>-, -CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>OH, -CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>NH<sub>2</sub>, -CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>Het', -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-Ar', -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-Het', -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-cycloalkyl, -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-COOA or -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>N(R<sup>4</sup>)<sub>2</sub>,

25

$R^4$  is H or A,

$W$  is N, CR<sup>3</sup> or an sp<sup>2</sup>-hybridised carbon atom,

$E$  together with  $W$  is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms,

30

which may contain a double bond,

35

$D$  is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or polysubstituted by Hal, A, OR<sup>3</sup>, N(R<sup>3</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>3</sup> or CON(R<sup>3</sup>)<sub>2</sub>,

- 
- 5            G        is  $-[C(R^4)_2]_n-$ ,  $-[C(R^4)_2]_nNR^3-$ ,  $-[C(R^4)_2]_nO-$ ,  $-[C(R^4)_2]_nS-$  or  
 $-[C(R^4)=C(R^4)]_n-$ ,
- 10          X        is  $-[C(R^4)_2]_nCONR^3[C(R^4)_2]_n-$ ,  $-[C(R^4)_2]_nNR^3CO[C(R^4)_2]_n-$ ,  
 $-[C(R^4)_2]_nNR^3[C(R^4)_2]_n-$ ,  $-[C(R^4)_2]_nO[C(R^4)_2]_n-$ ,  
 $-[C(R^4)_2]_nCO[C(R^4)_2]_n-$  or  $-[C(R^4)_2]_nCOO[C(R^4)_2]_n-$ ,
- 15          Y        is alkylene, cycloalkylene, Het-diyl or Ar-diyl,
- 20          T        is a monocyclic or bicyclic, saturated or unsaturated carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is monosubstituted or disubstituted by  $=O$ ,  $=S$ ,  $=NR^3$ ,  $=N-CN$ ,  $=N-NO_2$ ,  $=NOR^3$ ,  $=NCOR^3$ ,  $=NCOOR^3$  or  $=NOCOR^3$  and may furthermore be monosubstituted, disubstituted or trisubstituted by  $R^3$ , Hal, A,  $-[C(R^4)_2]_n-Ar$ ,  $-[C(R^4)_2]_n-Het$ ,  $-[C(R^4)_2]_n$ -cycloalkyl,  $OR^3$ ,  $N(R^3)_2$ ,  $NO_2$ , CN, COOR<sup>3</sup>, CON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>COA, NR<sup>3</sup>CON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>SO<sub>2</sub>A, COR<sup>3</sup>, SO<sub>2</sub>NR<sup>3</sup> and/or S(O)<sub>n</sub>A,
- 25          A        is unbranched or branched alkyl having 1-10 carbon atoms in which one or two CH<sub>2</sub> groups may be replaced by O or S atoms and/or by -CH=CH- groups and/or in addition 1-7 H atoms may be replaced by F,
- 30          Ar        is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR<sup>3</sup>, N(R<sup>3</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>3</sup>, CON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>COA, NR<sup>3</sup>CON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>SO<sub>2</sub>A, COR<sup>3</sup>, SO<sub>2</sub>N(R<sup>3</sup>)<sub>2</sub>, S(O)<sub>n</sub>A,  $-[C(R^4)_2]_n-COOR^3$  or  $-O[C(R^4)_2]_n-COOR^3$ ,
- 35          Ar'      is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR<sup>4</sup>, N(R<sup>4</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>4</sup>, CON(R<sup>4</sup>)<sub>2</sub>, NR<sup>4</sup>COA, NR<sup>4</sup>CON(R<sup>4</sup>)<sub>2</sub>, NR<sup>4</sup>SO<sub>2</sub>A, COR<sup>4</sup>, SO<sub>2</sub>N(R<sup>4</sup>)<sub>2</sub>, S(O)<sub>n</sub>A,  $-[C(R^4)_2]_n-COOR^4$  or  $-O[C(R^4)_2]_n-COOR^4$ ,
- Het     is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic ring having from 1 to 4 N, O and/or S atoms which may be unsubstituted or monosubstituted,

- disubstituted or trisubstituted by Hal, A,  $-[C(R^4)_2]_n-Ar$ ,  
-[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-Het', -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-cycloalkyl, OR<sup>3</sup>, N(R<sup>3</sup>)<sub>2</sub>,  
NR<sup>3</sup>CON(R<sup>3</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-COOR<sup>3</sup>,  
-[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-CON(R<sup>3</sup>)<sub>2</sub>, NR<sup>3</sup>COA, NR<sup>3</sup>SO<sub>2</sub>A, COR<sup>3</sup>, SO<sub>2</sub>NR<sup>3</sup>,  
S(O)<sub>m</sub>A and/or carbonyl oxygen,
- Het' is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic ring having from 1 to 4 N, O and/or S atoms which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, =S, =N(R<sup>4</sup>)<sub>2</sub>, Hal, A, OR<sup>4</sup>, N(R<sup>4</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>4</sup>, CON(R<sup>4</sup>)<sub>2</sub>, NR<sup>4</sup>COA, NR<sup>4</sup>CON(R<sup>4</sup>)<sub>2</sub>, NR<sup>4</sup>SO<sub>2</sub>A, COR<sup>4</sup>, SO<sub>2</sub>NR<sup>4</sup> and/or S(O)<sub>n</sub>A,
- Hal is F, Cl, Br or I,
- n is 0, 1 or 2,
- o is 1, 2 or 3;
- and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 20 2. Compounds according to Claim 1, in which  
D is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or disubstituted by Hal,  
25 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 30 3. Compounds according to Claim 1 or 2, in which  
D is phenyl, pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 35 4. Compounds according to one or more of Claims 1-3,  
in which

- R<sup>1</sup>, R<sup>2</sup> are each, independently of one another, H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, NHCOA, NHSO<sub>2</sub>A, OCH<sub>2</sub>COOA or OCH<sub>2</sub>COOH,  
5 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 10 5. Compounds according to one or more of Claims 1-4,  
in which  
G is (CH<sub>2</sub>)<sub>n</sub>, (CH<sub>2</sub>)<sub>n</sub>NH-, -CH=CH- or -CH=CH-CH=CH-,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 15 6. Compounds according to one or more of Claims 1-5,  
in which  
X is -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>CONR<sup>3</sup>[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
20
7. Compounds according to one or more of Claims 1-6,  
in which  
X is -CONH- or -CON(CH<sub>2</sub>COOA)-,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
25
- 30 8. Compounds according to one or more of Claims 1-7,  
in which  
Y is cycloalkylene, Het-diyl or Ar-diyl,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
35
9. Compounds according to Claims 1-8,

in which

Y is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene  
which is unsubstituted or monosubstituted or disubstituted by  
A, OA, Cl, F, COOCH<sub>3</sub>, COOH, phenoxy or aminocarbonyl,  
and pharmaceutically usable derivatives, solvates, salts and stereo-  
isomers thereof, including mixtures thereof in all ratios.

10. Compounds according to one or more of Claims 1-9,

in which

T is a monocyclic, saturated or unsaturated heterocyclic ring  
having 1 to 2 N and/or O atoms which is monosubstituted or  
disubstituted by =O, =S or =NH and may be monosubstituted  
or disubstituted by Hal, A and/or OA,  
and pharmaceutically usable derivatives, solvates, salts and stereo-  
isomers thereof, including mixtures thereof in all ratios.

20. 11. Compounds according to one or more of Claims 1-10,

in which

T is piperidin-1-yl, pyrrolidin-1-yl, pyridin-1-yl, morpholin-4-yl,  
piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl, pyrazin-1-yl,  
azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, imidazolidinyl,  
thiazolyl or 1,4-oxazepanyl, each of which is monosubstituted  
or disubstituted by =O or =NH and where the radicals may  
also be monosubstituted or disubstituted by Hal, A and/or  
OA,

30. and pharmaceutically usable derivatives, solvates, salts and stereo-  
isomers thereof, including mixtures thereof in all ratios.

35. 12. Compounds according to one or more of Claims 1-11,

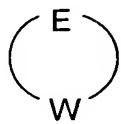
in which

- Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OA, SO<sub>2</sub>A, COOR<sup>2</sup>, SO<sub>2</sub>NH<sub>2</sub>, CN, COOA, COOH or phenoxy,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 5
13. Compounds according to one or more of Claims 1-12,  
in which
- 10 D is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or disubstituted by Hal,
- 15 R<sup>1</sup>, R<sup>2</sup> are each, independently of one another, H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, NHCOA, NHSO<sub>2</sub>A, OCH<sub>2</sub>COOA or OCH<sub>2</sub>COOH,
- 20 R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- R<sup>3</sup> is H, A, phenyl, benzyl or [C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>COOA,
- R<sup>4</sup> is H or A,
- 25 W is N, CR<sup>3</sup> or an sp<sup>2</sup>-hybridised carbon atom, E together with W is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms, which may contain a double bond,
- 30 G is (CH<sub>2</sub>)<sub>n</sub>, (CH<sub>2</sub>)<sub>n</sub>NH-, -CH=CH- or -CH=CH-CH=CH-, X is -[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>CONR<sup>3</sup>[C(R<sup>4</sup>)<sub>2</sub>]<sub>n</sub>-,
- Y is cycloalkylene, Het-diyl or Ar-diyl,
- 35 Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OA, SO<sub>2</sub>A, COOR<sup>2</sup>, SO<sub>2</sub>NH<sub>2</sub>, CN, COOA, COOH or phenoxy,

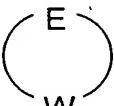
5           T       is a monocyclic, saturated or unsaturated heterocyclic ring having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by =O, =S or =NH and may be monosubstituted or disubstituted by Hal, A and/or OA,  
A       is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,  
Hal      is F, Cl, Br or I,  
n        is 0, 1 or 2,  
10       and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

14. Compounds according to one or more of Claims 1-13,  
15       in which  
D        is phenyl, pyridyl, thieryl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,  
R<sup>1</sup>, R<sup>2</sup>   are each, independently of one another, H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, NHCOA, NHSO<sub>2</sub>A, OCH<sub>2</sub>COOA or OCH<sub>2</sub>COOH,  
20       R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,  
R<sup>3</sup>      is H, A or CH<sub>2</sub>COOA,  
R<sup>4</sup>      is H or A,  
W        is N, CR<sup>3</sup> or an sp<sup>2</sup>-hybridised carbon atom,  
E        together with W is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms,  
30       which may contain a double bond,  
G        is (CH<sub>2</sub>)<sub>n</sub>, (CH<sub>2</sub>)<sub>n</sub>NH-, -CH=CH- or -CH=CH-CH=CH-,  
X        is -CONH- or -CON(CH<sub>2</sub>COOA)-,  
35       Y       is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene which is unsubstituted or monosubstituted or

- disubstituted by A, OA, Cl, F, COOCH<sub>3</sub>, COOH,  
phenoxy or aminocarbonyl,
- T      is piperidin-1-yl, pyrrolidin-1-yl, pyridin-1-yl, morpholin-4-  
5      yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl,  
      pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl,  
      imidazolidinyl, thiazolyl or 1,4-oxazepanyl, each of which  
      is monosubstituted or disubstituted by =O or =NH and  
      where the radicals may also be monosubstituted or  
10     disubstituted by Hal, A and/or OA,
- A      is unbranched or branched alkyl having 1-10 carbon  
      atoms and in which 1-7 H atoms may be replaced by F,  
Hal     is F, Cl, Br or I,  
n      is 0, 1 or 2,  
15     and pharmaceutically usable derivatives, solvates, salts and stereo-  
      isomers thereof, including mixtures thereof in all ratios.
15. Compounds according to one or more of Claims 1-14,  
20     in which
- D      is phenyl, pyridyl or thieryl, each of which is  
      monosubstituted or disubstituted by Hal,
- R<sup>1</sup>    is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4,  
25     5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy,  
      -OCOR<sup>3</sup>, NHCOA or NHSO<sub>2</sub>A,
- R<sup>2</sup>    is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 car-  
      bon atoms,
- 30     R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to  
      6-membered carbocyclic ring,
- R<sup>3</sup>    is H or A,
- R<sup>4</sup>    is H or A,



- is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,
- 5           G        is  $(CH_2)_n$  or  $(CH_2)_nNH-$ ,
- 10          X        is CONH,
- Y        is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,
- 15          T        is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo-[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen,
- 20          A        is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- Hal      is F, Cl, Br or I,
- n        is 0, 1 or 2;
- 25          and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
16. Compounds according to one or more of Claims 1-15,  
30          in which
- D        is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,
- R<sup>1</sup>    is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, -OCOR<sup>3</sup>, NHCOA or NHSO<sub>2</sub>A,
- 35

$R^2$  is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,  
 $R^1$  and  $R^2$  together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,  
5       $R^3$  is H or A,  
 $R^4$  is H or A,  
 is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-  
10     3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,  
15     G is  $(CH_2)_n$  or  $(CH_2)_nNH-$ ,  
X is CONH,  
Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,  
20     T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,  
A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F;  
25     Hal is F, Cl, Br or I,  
n is 0, 1 or 2;  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
30

17. Compounds according to one or more of Claims 1-16,  
in which

35     X is  $-[C(R^4)_2]_nCONR^3[C(R^4)_2]_n-$  or  $-[C(R^4)_2]_nCO[C(R^4)_2]_n-$ ,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

18. Compounds according to one or more of Claims 1-17,

in which

X is CONH or COCH<sub>2</sub>,

5 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

19. Compounds according to one or more of Claims 1-18,

10 in which

D is phenyl, pyridyl or thiienyl, each of which is monosubstituted or disubstituted by Hal,

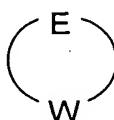
R<sup>1</sup> is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 15 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, -OCOR<sup>3</sup>, NHCOA or NHSO<sub>2</sub>A,

R<sup>2</sup> is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

20 R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,

R<sup>3</sup> is H or A,

R<sup>4</sup> is H or A,

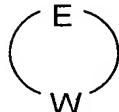
25  is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-

3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or 30 azetidine-1,2-diyl,

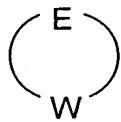
G is (CH<sub>2</sub>)<sub>n</sub> or (CH<sub>2</sub>)<sub>n</sub>NH<sub>2</sub>,

X is CONH or COCH<sub>2</sub>,

35 Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

- T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,
- A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,  
5 Hal is F, Cl, Br or I,  
n is 0, 1 or 2,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 10
20. Compounds according to one or more of Claims 1-19, in which
- D is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,
- 15 R<sup>1</sup> is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, -OCOR<sup>3</sup>, NHCOA, NHSO<sub>2</sub>A, H-C≡C-CH<sub>2</sub>-, CH<sub>3</sub>-C≡C-CH<sub>2</sub>-O-, -O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>OH, -O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>NH<sub>2</sub> or -O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>Het',  
20 R<sup>2</sup> is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,
- R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,  
25 R<sup>3</sup> is H or A,  
R<sup>4</sup> is H or A,  
30  is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,  
35 G is (CH<sub>2</sub>)<sub>n</sub> or (CH<sub>2</sub>)<sub>n</sub>NH-,

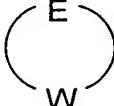
- X is CONH or COCH<sub>2</sub>,
- Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,
- 5 T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,
- Het' is a saturated 3-6-membered heterocyclic ring having from 1 to 3 N and/or O atoms, which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, Hal, A, OH, NH<sub>2</sub>, NO<sub>2</sub>, CN, COOA or CONH<sub>2</sub>,
- 10 A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- 15 Hal is F, Cl, Br or I,
- n is 0, 1 or 2,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 20 21. Compounds according to one or more of Claims 1-20,  
in which
- D is phenyl, pyridyl or thienyl, each of which is mono-substituted or disubstituted by Hal,
- 25 R<sup>1</sup> is ethynyl, vinyl, allyloxy, CH<sub>3</sub>-C≡C-CH<sub>2</sub>-O-,  
-O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>OH, -O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>NH<sub>2</sub> or  
-O-CH<sub>2</sub>-CH(OH)-CH<sub>2</sub>Het',
- R<sup>2</sup> is H or OH,
- 30 R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- R<sup>3</sup> is H or A,
- R<sup>4</sup> is H or A,



- is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,
- 5           G        is  $(CH_2)_n$  or  $(CH_2)_nNH-$ ,
- 10          X        is CONH, CO, COO or COCH<sub>2</sub>,
- Y        is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,
- 15          T        is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo-[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen or OA,
- 20          Het'     is a saturated 3-6-membered heterocyclic ring having from 1 to 3 N and/or O atoms, which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, Hal, A, OH, NH<sub>2</sub>, NO<sub>2</sub>, CN, COOA or CONH<sub>2</sub>,
- 25          A        is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- Hal      is F, Cl, Br or I,
- n        is 0, 1 or 2,
- 30          and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

22. Compounds according to one or more of Claims 1-21,  
in which

35          D        is phenyl, pyridyl, thiophenyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,

|    |   |   |
|----|---|---|
|    | $R^1$   | is H, =O, COOR <sup>3</sup> , OH, OA, NH <sub>2</sub> , alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N <sub>3</sub> , ethynyl, vinyl, allyloxy, NHCOA, NHSO <sub>2</sub> A, OCH <sub>2</sub> COOA or OCH <sub>2</sub> COOH,                                      |
| 5  | $R^2$   | is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,   |
|    | $R^1$ and $R^2$   | together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,  |
| 10 | $R^3$   | is H or A,  |
|    | $R^4$   | is H or A,  |
|    |  | is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1 <i>H</i> -pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl, |
| 15 | $G$   | is (CH <sub>2</sub> ) <sub>n</sub> , (CH <sub>2</sub> ) <sub>n</sub> NH-, -CH=CH- or -CH=CH-CH=CH-,   |
| 20 | $X$   | is CONH, COCH <sub>2</sub> or -CON(CH <sub>2</sub> COOA)-,  |
|    | $Y$   | is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene which is unsubstituted or monosubstituted or disubstituted by A, OA, Cl, F, COOCH <sub>3</sub> , COOH, phenoxy or aminocarbonyl,   |
| 25 | $T$   | is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,   |
|    | $A$   | is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,   |
| 30 | Hal   | is F, Cl, Br or I,  |
|    | $n$   | is 0, 1 or 2,   |
|    |   | and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.   |
| 35 |   |   |

23. Compounds according to Claim 1, selected from the group consisting of

- 5            1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-  
             (R)-pyrrolidine-1,2-dicarboxamide,
- 10          1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-  
             phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,
- 15          1-N-[(4-chlorophenyl)]-2-N-{{[3-fluoro-4-(3-oxomorpholin-4-yl)-  
             phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,
- 20          1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
             phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,
- 25          1-N-[(4-chlorophenyl)]-2-N-{{[3-trifluoromethyl-4-(3-oxomorpho-  
             lin-4-yl)phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,
- 30          1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-  
             phenyl]}-(R)-piperidine-1,2-dicarboxamide,
- 35          1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-  
             (R)-pyrrolidine-1,2-dicarboxamide,
- 1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-  
             (R)-pyrrolidine-1,2-dicarboxamide,
- 1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-  
             (R)-2,5-dihydropyrrole-1,2-dicarboxamide,
- N-[4-(3-oxomorpholin-4-yl)phenyl]-(R)-1-(5-chlorothiophene-  
             2-carbonyl)pyrrolidine-2-carboxamide,
- N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-(R)-1-(5-chloro-  
             thiophene-2-carbonyl)pyrrolidine-2-carboxamide,
- 3-N-[(4-chlorophenyl)]-4-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-  
             (R)-oxazolidine-3,4-dicarboxamide,
- 3-N-[(4-chlorophenyl)]-4-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-  
             phenyl]}-(R)-oxazolidine-3,4-dicarboxamide,
- 3-N-[(4-chlorophenyl)]-4-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-  
             (4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

- 3-N-[(4-chlorophenyl)-4-N-{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,  
5 3-N-[(4-chlorophenyl)]-4-N-{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(R)-oxazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,  
10 3-N-[(4-chlorophenyl)]-4-N-{[3-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,  
15 3-N-[(4-chlorophenyl)]-4-N-{[3-chloro-4-(3-oxomorpholin-4-yl)-phenyl]}-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(4R,5R)-5-methyloxazolidine-3,4-dicarboxamide,  
20 3-N-[(4-chlorophenyl)]-4-N-{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(R)-oxazolidine-3,4-dicarboxamide,  
25 3-N-[(4-chlorophenyl)]-4-N-{[3-chloro-4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(R)-oxazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(S)-thiazolidine-3,4-dicarboxamide,  
30 3-N-[(4-chlorophenyl)]-4-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(S)-1,1-dioxo-1 $\lambda^6$ -thiazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(S)-thiazolidine-3,4-dicarboxamide,  
35 3-N-[(4-chlorophenyl)]-4-N-{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(S)-1,1-dioxo-1 $\lambda^6$ -thiazolidine-3,4-dicarboxamide,  
3-N-[(4-chlorophenyl)]-4-N-{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(R)-thiazolidine-3,4-dicarboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,  
N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,

- N-[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,  
1-N-[(5-chloropyridin-2-yl)]-2-N-{{[4-(2-oxo-2*H*-pyridin-1-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
5 1-N-[(5-chloropyridin-2-yl)]-2-N-{{[4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(5-chloropyridin-2-yl)]-2-N-{{[4-(2-oxo-2*H*-pyrazin-1-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
10 1-N-[(5-chloropyridin-2-yl)]-2-N-{{[3-fluoro-4-(2-oxo-2*H*-pyridin-1-  
yl)phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(5-chloropyridin-2-yl)]-2-N-{{[4-(2-oxo-2*H*-pyridin-1-yl)-  
phenyl]}-(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,  
15 1-N-[(5-chloropyridin-2-yl)]-2-N-{{[4-(3-oxomorpholin-4-yl)-  
phenyl]}-(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]}-  
(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]}-  
(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxopyrazin-1-yl)phenyl]}-  
(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[3-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)-  
phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
35 1-N-[(4-chlorophenyl)]-2-N-{{[3-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2*R*,3*R*)-3-hydroxypyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-{[3-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,3S)-3-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
5 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2S,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-3,4-dihydroxypyrrolidine-1,2-dicarboxamide,  
10 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-azidopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-aminopyrrolidine-1,2-dicarboxamide,  
15 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-azidopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-aminopyrrolidine-1,2-dicarboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-acetaminopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-acetaminopyrrolidine-1,2-dicarboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-methylsulfonylaminopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-methylsulfonylaminopyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
35 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-propoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-allyloxypprrolidine-1,2-dicarboxamide,

- (3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl isobutyrate,  
5 (3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl propionate,  
(3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl acetate,  
10 4-N-[(4-chlorophenyl)]-5-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-1,3-dioxolane-4,5-dicarboxamide,  
4-N-[(4-chlorophenyl)]-5-N-{[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]}-1,3-dioxolane-4,5-dicarboxamide,  
15 4-N-[(4-chlorophenyl)]-5-N-{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-1,3-dioxolane-4,5-dicarboxamide,  
4-N-[(4-chlorophenyl)]-5-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,  
20 4-N-[(4-chlorophenyl)]-5-N-{[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]}-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,  
4-N-[(4-chlorophenyl)]-5-N-{[4-(2-oxo-1H-pyridin-1-yl)phenyl]}-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,  
25 1-N-[4-chlorophenyl]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-1-BOC-piperazine-1,2-dicarboxamide,  
1-N-[4-chlorophenyl]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-piperazine-1,2-dicarboxamide,  
30 1-N-[4-chlorophenyl]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-1,3-oxazinane-3,4-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,  
35 6-N-[(4-chlorophenyl)]-7-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-4-oxa-6-azaspiro[2.4]heptane-6,7-dicarboxamide,  
1-N-[(6-chloropyridin-3-yl)]-2-N-{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(6-chloropyridin-3-yl)]-2-N-{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4S)-4-acetaminopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-butylsulfonylaminopyrrolidine-1,2-dicarboxamide,  
5 1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-(R)-4-oxopyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4S)-4-aminopyrrolidine-1,2-dicarboxamide,  
10 1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(S)-pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-hydroxypyrrrolidine-1,2-dicarboxamide,  
15 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[2-(4-chloro-phenyl)acetyl]-4-hydroxypyrrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(4-chlorobenzoyl)-4-hydroxypyrrrolidine-2-carboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-methoxypyrrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-(2R,4R)-4-methoxypyrrrolidine-1,2-dicarboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-methoxypyrrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(2R,4R)-4-methoxypyrrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4S)-4-(2-methylpropanoylamino)pyrrolidine-1,2-dicarboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(1-1*H*-indol-3-yl-methanoyl)-4-hydroxypyrrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(1-1*H*-indol-6-yl-methanoyl)-4-hydroxypyrrrolidine-2-carboxamide,  
35

- 1-N-[(4-chlorophenyl)]-2-N-{{[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-1H-pyridin-1-yl)phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
5 1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-1H-pyridin-1-yl)phenyl]}-(2R,4S)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,  
10 1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(2R,4S)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-2H-pyridin-1-yl)phenyl]}-4,4-difluoro-(R)-pyrrolidine-1,2-dicarboxamide,  
15 1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(2-oxo-2H-pyridin-1-yl)-phenyl]}-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(2-oxo-2H-pyridin-1-yl)-phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{{[2-fluoro-4-(2-oxo-2H-pyridin-1-yl)-phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
2-N-[(4-chlorophenyl)]-1-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,  
25 2-N-[(4-chlorophenyl)]-1-N-{{[4-(3-oxomorpholin-4-yl)phenyl]}-(S)-pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-3-methoxy-2H-pyridin-1-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{{[4-(2-oxo-3-methoxy-2H-pyridin-1-yl)phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,  
N-(4-chlorophenyl)-(R)-1-{2-[4-(3-oxomorpholin-4-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(S)-1-{2-[4-(3-oxomorpholin-4-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,  
35

- N-(4-chlorophenyl)-(2R,4R)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(2R,4S)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,  
5 N-(4-chlorophenyl)-(2S,4R)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(S)-1-{2-[4-(2-oxo-1H-pyridin-1-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,  
10 N-(4-chlorophenyl)-(S)-1-{2-[4-(2-oxopyrrolidin-1-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(R)-1-{2-[4-(2-oxopyrrolidin-1-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,  
15 N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)phenyloxy-carbonyl]pyrrolidine-2-carboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{[4-(2-oxo-2H-pyrazin-1-yl)phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[3-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(prop-2-ynyoxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(but-2-ynyoxy)pyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(2,3-dihydroxypropoxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)pyrrolidine-1,2-dicarboxamide,  
35 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(2-oxooxazolidin-5-ylmethoxy)pyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2R,4R)-4-(3-amino-2-hydroxypropoxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(2-oxo-1*H*-pyrazin-1-yl)phenyl]}-  
(R)-2,5-dihydropyrrole-1,2-dicarboxamide,  
5 1-N-[(4-chlorophenyl)]-2-N-{[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]}-  
(R)-2,5-dihydropyrrole-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[3-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(R)-2,5-dihydropyrrole-1,2-dicarboxamide,  
10 1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(R)-2,5-dihydropyrrole-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2S,3S)-3-hydroxypyrrolidine-1,2-dicarboxamide,  
15 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2S,4S)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[2-methoxycarbonyl-4-(3-oxomor-  
pholin-4-yl)phenyl]}-(2R,4R)-3-hydroxypyrrolidine-1,2-dicarboxamide,  
20 1-N-[(4-chlorophenyl)]-2-N-{[2-carboxy-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2R,4R)-3-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2R,3S,4R)-3,4-dihydroxypyrrolidine-1,2-dicarboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2R,4R)-4-allyloxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2R,4R)-4-(prop-2-ynloxy)pyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2R,4S)-4-(prop-2-ynloxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2R,4R)-4-(methoxycarbonylmethoxy)pyrrolidine-1,2-dicarboxamide,  
35 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-  
(2R,4R)-4-(carboxymethoxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-bromophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-  
phenyl]}-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]}-(2R,4R)-4-(2,3-dihydroxypropoxy)pyrrolidine-1,2-dicarboxamide,
- 5 1-N-[(4-chlorophenyl)]-2-N-{N-methoxycarbonylmethyl-N'-[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,
- 10 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)cyclohexan-1-yl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,
- 15 1-N-[(4-chlorophenyl)]-2-N-{[4-(2-iminopyrrolidin-1-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 442;
- 20 1-N-[(4-chlorophenyl)]-2-N-{[3-methyl-4-(2-iminopyrrolidin-1-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 456;
- 25 1-N-[(4-chlorophenyl)]-2-N-[4-{2-[(E)-cyanimino]imidazolidin-1-yl}phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 468;
- 30 1-N-[(4-chlorophenyl)]-2-N-{[4-(2-imino-5-methylthiazol-3-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 473;
- 35 1-N-[(4-chlorophenyl)]-2-N-{[2-aminocarbonyl-4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 502;
- 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxy-2-methylpyrrolidine-1,2-dicarboxamide,
- 25 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,
- 30 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-thiophen-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,
- 35 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(2E,4E)-5-phenylpenta-2,4-dienyloyl]-4-hydroxypyrrolidine-2-carboxamide,
- N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-methylfuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,
- N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-thiophen-2-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

- N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
5 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
10 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
15 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
20 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
25 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
30 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
35 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

- N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(5-chlorofuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
5 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(5-chlorofuran-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(4-chlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
10 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(3,4-dichlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(4-chlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
15 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(5-chlorofuran-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-(5-chlorothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
20 N-[4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-1*H*-imidazol-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-1*H*-imidazol-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
25 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-1*H*-imidazol-4-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,  
N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-1*H*-imidazol-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
30 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-pyridin-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-pyridin-3-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
35 N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]- $(2R,4R)$ -1-[ $(E)$ -3-pyridin-3-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,

- N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
5 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,  
10 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,  
15 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-bromo-thiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-bromo-thiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
20 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-bromo-thiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,  
N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-bromo-thiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,  
25 N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,  
N-(4-chlorophenyl)-(S)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-{[4-(5-oxo-1,4-oxazepan-4-yl)-phenyl]}-(R)-pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(5-oxo-1,4-oxazepan-4-yl)-phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
35 1-N-[(4-chlorophenyl)]-2-N-{[4-((S)-2-methyl-3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-((S)-2-methyl-3-oxomorpholin-4-yl)phenyl]}-(2R)-pyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-{[4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]}-(2R)-pyrrolidine-1,2-dicarboxamide,  
5 1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)-2-phenoxyphenyl]}-(2R)-pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[2-fluoro-4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
10  
15 1-N-[(4-chlorophenyl)]-3-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-piperidine-1,3-dicarboxamide,  
1-N-[(4-chlorophenyl)]-3-N-{[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]}piperidine-1,3-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-(2-methoxyethoxy)pyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxo-1,4-oxazepan-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
20  
25 1-N-[(4-chlorophenyl)]-2-N-{[2-methyl-4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-{[2-(3-oxomorpholin-4-yl)phenyl]}-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
30  
35 24. Pyrrolidinecarboxylic acid derivatives selected from the group consisting of  
1-N-[(4-chlorophenyl)]-2-N-[(1'-methyl-[1,4']bipiperidinyl-4-yl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

- 1-N-[(4-chlorophenyl)]-2-N-[(3,4,5,6-tetrahydro-2H-1,4'-bipyridinyl-4-yl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-[(3,4,5,6-tetrahydro-2H-1,4'-bipyridinyl-4-yl)]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,  
5 N-(4-chlorophenyl)-(2R,4R)-4-hydroxy-2-(4-pyridin-4-yl)piperazine-1-carbonyl)pyrrolidine-1-carboxamide,  
N-(4-chlorophenyl)-(2R,4R)-4-hydroxy-2-[4-(2-methoxyphenyl)-piperazine-1-carbonyl]pyrrolidine-1-carboxamide,  
10 N-(4-chlorophenyl)-(2R,4R)-2-[4-(4-fluorophenyl)piperazine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,  
N-(4-chlorophenyl)-(2R,4R)-4-hydroxy-2-[4-hydroxy-4-(4-methoxyphenyl)piperidine-1-carbonyl]pyrrolidine-1-carboxamide,  
15 N-(4-chlorophenyl)-(2R,4R)-4-hydroxy-2-(4-pyridin-2-yl)piperazine-1-carbonyl)pyrrolidine-1-carboxamide,  
N-(4-chlorophenyl)-(2R,4R)-2-[4-(4-ethylpiperazin-1-yl)piperidine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,  
20 N-(4-chlorophenyl)-(2R,4R)-2-[4-(4,6-dimethylpyrimidin-2-yl)-piperazine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,  
N-(4-chlorophenyl)-(2R,4R)-4-hydroxy-2-[4-(1-methylpiperidin-4-yl)piperazine-1-carbonyl]pyrrolidine-1-carboxamide,  
25 1-N-[(4-chlorophenyl)]-2-N-[(2-(2-dimethylaminoethoxy)-4-morpholin-4-ylphenyl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
1-N-[(4-chlorophenyl)]-2-N-[(2-ethoxy-4-morpholin-4-ylphenyl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
30 1-N-[(4-chlorophenyl)]-2-N-[(4-morpholin-4-yl-2-propoxyphenyl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,  
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.  
35 25. Cyclopentanecarboxylic acid derivatives selected from the group consisting of

N-[4-(3-oxomorpholin-4-yl)phenyl]-(rac)-2-[3-(4-chlorophenyl)-ureido]cyclopentanecarboxamide,

5 N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-(rac)-2-[3-(4-chlorophenyl)ureido]cyclopentanecarboxamide,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

10

26. Process for the preparation of compounds of the formula I according to Claims 1-23 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, characterised in that

15

a) for the preparation of compounds of the formula I in which

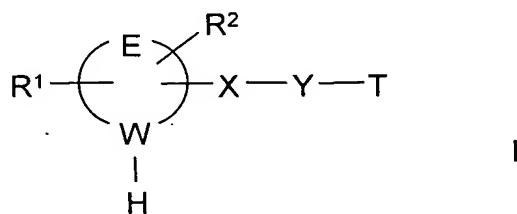
W is N and

G is NH,

20

a compound of the formula II

25



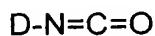
in which

R<sup>1</sup>, R<sup>2</sup>, E, X, Y and T are as defined in Claim 1,  
and W is N,

30

is reacted with a compound of the formula III

35



III

in which

D is as defined in Claim 1,

or

5

- b) for the preparation of compounds of the formula I in which  
 $X$  is  $-[C(R^4)_2]_nCONR^3[C(R^4)_2]_n-$ ,

a compound of the formula IV

10

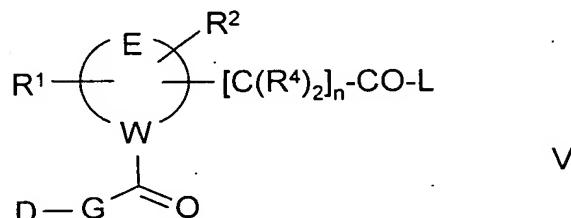


in which  $R^3$ ,  $n$ ,  $Y$  and  $T$  are as defined in Claim 1,

15

is reacted with a compound of the formula V

20



in which

25

$L$  is Cl, Br, I or a free or reactively functionally modified OH group,  
 and

$R^1$ ,  $R^2$ ,  $R^4$ ,  $D$ ,  $E$ ,  $G$ ,  $W$  and  $n$  are as defined in Claim 1,

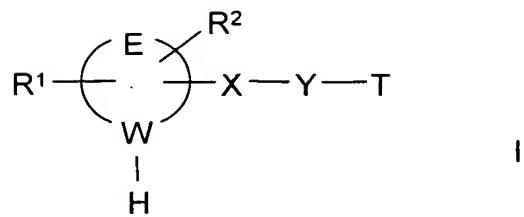
30

or

- c) for the preparation of compounds of the formula I in which  $W$  is N,

35

a compound of the formula II



5

in which

$R^1$ ,  $R^2$ ,  $E$ ,  $X$ ,  $Y$  and  $T$  are as defined in Claim 1, and  $W$  is  $N$ .

10

is reacted with a compound of the formula VI



15

in which D and G are as defined in Claim 1, and

L is Cl, Br, I or a free or reactively functionally modified OH group,

2

and/or

20

a base or acid of the formula I is converted into one of its salts.

25

27. Compounds of the formula I according to one or more of Claims 1 to 23 and the compounds of Claims 24 and 25 as inhibitors of coagulation factor Xa

30

28. Compounds of the formula I according to one or more of Claims 1 to 23 and the compounds of Claims 24 and 25 as inhibitors of coagulation factor VIIa.

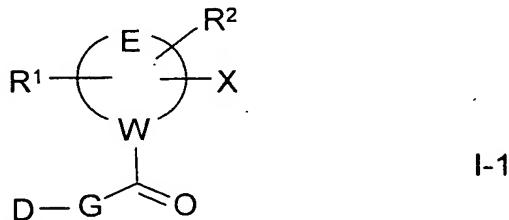
29. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, and, if desired, excipients and/or adjuvants.

30. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and at 5 least one further medicament active ingredient.
10. 31. Use of compounds according to one or more of Claims 1 to 23 or the compounds of Claims 24 and 25 and/or physiologically acceptable salts, salts and solvates thereof for the preparation of a medicament for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis 15 after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases.
20. 32. Set (kit) consisting of separate packs of  
(a) an effective amount of a compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, and
25. (b) an effective amount of a further medicament active ingredient.
30. 33. Use of compounds of the formula I according to one or more of Claims 1 to 23 or of compounds of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis 35 after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases,

in combination with at least one further medicament active ingredient.

34. Intermediate compounds of the formula I-1

5



10

in which

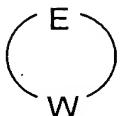
D is phenyl, pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,

R<sup>1</sup> is H, OH, OA, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms or ethynyl,

15

R<sup>2</sup> is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

20

 is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl,

G is (CH<sub>2</sub>)<sub>n</sub>, (CH<sub>2</sub>)<sub>n</sub>NH-, -CH=CH- or -CH=CH-CH=CH-,

X is COOH,

25

A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

Hal is F, Cl, Br or I,

n is 0, 1 or 2,

and isomers and salts thereof.

30

35. Compounds according to Claim 34, selected from the group consisting of

3-(4-chlorophenylcarbamoyl)oxazolidine-4-carboxylic acid,

3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxylic acid,

35

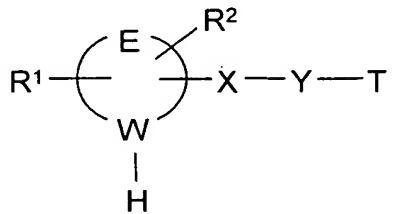
and isomers and salts thereof.

36. Intermediate compounds selected from the group consisting of  
 (2R,4S)-BOC-4-ethynyl-4-hydroxy-pyrrolidine-2-carboxylic acid,  
 (2R,4R)-BOC-4-ethynyl-4-hydroxy-pyrrolidine-2-carboxylic acid,  
 alkyl (2R,4S)-BOC-4-ethynyl-4-hydroxypyrrolidine-2-carboxylate,  
 alkyl (2R,4R)-BOC-4-ethynyl-4-hydroxypyrrolidine-2-carboxylate,  
 where alkyl has 1, 2, 3, 4, 5 or 6 carbon atoms,  
 and isomers and salts thereof.

5

10 37. Intermediate compounds of the formula I-2

15



I-2

in which

20

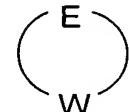
R<sup>1</sup> is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, NHCOA, NHSO<sub>2</sub>A, OCH<sub>2</sub>COOA or OCH<sub>2</sub>COOH,

25 R<sup>2</sup> is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

25

R<sup>1</sup> and R<sup>2</sup> together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,

30 R<sup>3</sup> is H or A,



30 is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-

3,4- or 3,5-diyl,

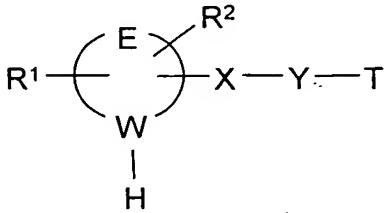
X is CONH,

35

Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

T        is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or  
 5        2-azabicyclo[2.2.2]octan-2-yl, each of which is mono-substituted or disubstituted by carbonyl oxygen,  
 A        is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,  
 Hal      is F, Cl, Br or I,  
 n        is 0, 1 or 2,  
 10        and isomers and salts thereof.

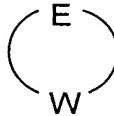
## 38. Compounds according to Claim 37 of the formula I-2a

15         I-2a

20        in which  
 R<sup>1</sup>      is H, =O, COOR<sup>3</sup>, OH, OA, NH<sub>2</sub>, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N<sub>3</sub>, ethynyl, vinyl, allyloxy, NHCOA, NSO<sub>2</sub>A, OCH<sub>2</sub>COOA or OCH<sub>2</sub>COOH,

25        R<sup>2</sup>      is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

R<sup>3</sup>      is H or A,

30        

X        is CONH,

Y        is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

T is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo-[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen,  
5  
A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,  
Hal is F, Cl, Br or I,  
n is 0, 1 or 2,  
10 and isomers and salts thereof.

39. Compounds according to Claim 38, selected from the group consisting of  
15      *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(S)-pyrrolidine-2-carboxamide,  
          *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(R)-pyrrolidine-2-carboxamide,  
          *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(2*R*,4*R*)-4-hydroxypyrrolidine-2-  
          carboxamide,  
20      *N*-[4-(3-oxomorpholin-4-yl)phenyl]-4-hydroxypyrrolidine-2-carbox-  
          amide,  
          *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(R)-4,4-dimethoxypyrrolidine-2-  
          carboxamide,  
25      *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(2*R*,4*R*)-4-methoxypyrrolidine-2-  
          carboxamide,  
          and isomers and salts thereof.
40. Medicament according to Claim 30, comprising 1-*N*-[(4-chloro-  
30      phenyl)]-2-*N*-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2*R*,4*R*)-4-hydroxy-  
          pyrrolidine-1,2-dicarboxamide and/or pharmaceutically usable  
          derivatives, solvates, salts and stereoisomers thereof, including  
          mixtures thereof in all ratios, and aspirin.

- 154 -

41. Use according to Claim 33, comprising 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, in combination with aspirin.

5

10

15

20

25

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